

銅存在下 6 π -電子環状反応を用いたピリド [4,3-*b*] カルバゾールアルカロイドの簡便な全合成

伊藤智貴^{*}、阿部 匠^{*}、町支臣成、西山卓志、柳田玲子^{**}、石倉 稔^{*}

Eur. J. Org. Chem., 2290-2299 (2016)

Concise Total Synthesis of Pyrido[4,3-*b*]carbazole Alkaloids Using Copper-Mediated 6 π -Electrocyclization

Tomoki Itoh^{*}, Takumi Abe^{*}, Tominari Choshi, Takashi Nishiyama,
Reiko Yanada^{**}, and Minoru Ishikura^{*}

ABSTRACT: Concise syntheses of 9-methoxyellipticine, 3,4-dihydroellipticine (μ -alkaloid D), 1,2,3,4-tetrahydroellipticine, 2-methyl-1,2,3,4-tetrahydroellipticine, olivacine, 3,4-dihydroolivacine, (\pm)-guatambuine, and (\pm)-janetine were developed starting from hexatriene intermediates readily obtained by Pd-catalyzed tandem cyclization/cross-coupling reaction of indolylborates. The route enables the facile construction of pyrido[4,3-*b*]carbazoles by Cu-catalyzed 6 π -electrocyclization and subsequent transformation of the pyridocarbazole intermediates into pyrido[4,3-*b*]carbazole alkaloids.

抄録 インドリルボラートの Pd 触媒によるタンデム環化 / クロスカップリング反応によって容易に得られるヘキサトリエン中間体を經由した 9-methoxyellipticine、3,4-dihydroellipticine(μ -alkaloid D)、1,2,3,4-tetrahydroellipticine、2-methyl-1,2,3,4-tetrahydroellipticine、olivacine、3,4-dihydroolivacine、(\pm)-guatambuine および (\pm)-janetine の簡便な合成法を開発した内容である。

^{*} School of Pharmaceutical Sciences, Health Science University of Hokkaido
北海道医療大学薬学部

^{**} Faculty of Pharmaceutical Sciences, Hiroshima International University
広島国際大学薬学部